

Original paper

CT-measured pulmonary artery diameter as an independent predictor of pulmonary hypertension in cystic fibrosis

Mehrdad Bakhshayeshkaram^{1A,D,E}, Farahnaz Aghahosseini^{1A,B,C,D,E,F}, Fatemeh Vaezi^{1B}, Shahram Kahkoei^{1D,E},
Yalda Salehi^{2E}, Maryam Hassanzad^{1A,B,C,D,E,F}, Hamid Reza Jamaati^{3D,E,F}, Ali Akbar Velayati^{4A,D,F}

¹Pediatric Respiratory Diseases Research Center, National Research Institute of Tuberculosis and Lung Diseases (NRITLD), Shahid Beheshti University of Medical Sciences, Tehran, Iran

²Imam Khomeini Hospital, Tehran University of Medical Science, Tehran, Iran

³Chronic Respiratory Diseases Research Center, National Research Institute of Tuberculosis and Lung Diseases (NRITLD), Shahid Beheshti University of Medical Sciences, Tehran, Iran

⁴Mycobacteriology Research Center (MRC), National Research Institute of Tuberculosis and Lung Diseases (NRITLD), Shahid Beheshti University of Medical Sciences, Tehran, Iran

Abstract

Purpose: The role of computed tomography (CT) scan, as a promising prognostic imaging modality in cystic fibrosis (CF), has been widely investigated, focusing on parenchymal abnormalities. The aim of the present study was to evaluate the diagnostic performance of thoracic vascular parameters on CT to detect pulmonary hypertension (PH).

Material and methods: CF patients who contemporaneously underwent CT and echocardiography were retrospectively enrolled. Baseline characteristics in addition to pulmonary artery diameter (PAD) and pulmonary to aortic (PA/A) ratio were compared between cohorts with and without PH, based on the results of echocardiography separately in paediatric patients (< 18) and adults (≥ 18).

Results: Of a total 119 CF patients, 39 (32.8%) had PH (paediatric: 23/78, 29.5%, adult: 16/41, 39%). In paediatric CF patients, mean age, HCO₃, PCO₂, and pulmonary artery diameter (PAD) were significantly higher in the PH group compared to the non-PH group. Mean pulmo however, only PAD remained as the independent predictor of PH based on multivariate analysis (overall: 22.86 mm [±3.86] vs. 18.43 mm [±4.72], $p = 0.005$, paediatric patients: 22.63 mm [±4.4] vs. 17.10 mm [±4.64], $p = 0.03$). Using a cut off of 19.25 mm, the diagnostic performance of PAD to detect PH was found to be as follows: sensitivity = 82%, specificity = 70%, and accuracy = 73.1%. No significant difference was demonstrated in PAD between PH and non-PH groups in adults with CF (23.19 [±3.60] vs. 21.34 [±3.49], $p = 0.7$).

Conclusions: In CF patients, PAD revealed an age-dependent performance to detect PH. PAD can be applied to predict pulmonary hypertension in paediatric CF patients and may be recommended to be routinely measured on follow-up chest CT scan in childhood CF.

Key words: cystic fibrosis, pulmonary hypertension, CT scan, pulmonary artery diameter, pulmonary artery diameter in CF pulmonary pressure.

Introduction

Pulmonary hypertension (PH) is commonly encountered in advanced cystic fibrosis (CF) and is associated with

a poor prognosis. The results of a recent study demonstrated that, regardless of the severity, the hazard ratio of pulmonary hypertension and related impact on mortality is significantly higher in CF patients than that of other

Correspondence address:

Assoc. Prof. Maryam Hassanzad, Pediatric Respiratory Diseases Research Center, National Research Institute of Tuberculosis and Lung Diseases (NRITLD), Massih Daneshvari Hospital Daar-Abad, Niavaran Ave., Tehran, Iran, zip code: 19569-44413, post box: 19575-154, phone: 021-27122712, e-mail: Mar_hassanzad@yahoo.com

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lung diseases [1]. Prompt in a multidisciplinary therapeutic approach, including drug treatment, nutritional support and respiratory rehabilitation, early detection of PH in CF patients may hence contribute to prevent or slow disease progression, improving quality of life and probably survival. Echocardiography, as the most sensitive non-invasive screening tool for PH, may not perform as a reliable tool in advanced lung parenchymal diseases, which frequently occurred in CF patients [2,3]. In addition, pulmonary function test may not correlate well with the course and severity of pulmonary parenchymal and vascular disease [4,5]. Right-sided heart catheterisation as the gold standard for diagnosis of PH with the capability of providing additional valuable information for patient stratification and treatment strategies is not frequently indicated in CF patients and may be associated with some major complications [6]. There is an increasing body of evidence suggesting a potential role for CT-measured vascular parameters to identify pulmonary hypertension. The results of a recent meta-analysis on 2134 cases from 20 publications revealed that pulmonary artery diameter and pulmonary-to-aorta ratio had a modest sensitivity (79%) and specificity (81%) with a good overall accuracy to detect pulmonary hypertension and may play a complementary role in the diagnostic approach to PH [7]. Providing details about both vascular and parenchymal components, CT may be considered as the imaging modality of choice for disease monitoring in progressive lung diseases; however, evidence about the diagnostic and prognostic implication of CT vascular metric in CF patients is scarce. A recent study revealed that the PA/A ratio has a strong correlation with acute respiratory exacerbation and also with cystic fibrosis transmembrane conductance regulator (CFTR) dysfunction in adults with CF and hence may provide valuable prognostic information in this cohort [8]. In addition, few studies are available investigating normal values of thoracic vasculature in the paediatric population [9] as well as any relation between vascular parameters and PH. The aim of the present study was to investigate the relationship between chest CT vascular parameters and pulmonary artery systolic pressure in CF patients, as well as their performance characteristics to detect PH, separately in paediatric and adult groups.

Material and methods

Patients

In this retrospective chart review study, all patients with proven diagnosis of CF admitted to the paediatric wards of Masih Daneshvari Hospital between 2012 and 2016 were consecutively included. Patients contemporaneously investigated by CT scan and echocardiography with a maximum interval of two weeks were then selected. Cardiopulmonary co-morbidity and PH with aetiologies

other than CF and aortic aneurysm/dilatation were considered as exclusion criteria. The study was approved by the Institutional Review Board, and the need for informed consent was waived. Demographic and anthropometric characteristics including age, sex, body mass index (BMI), and arterial blood gas indices including pH, PCO_2 , HCO_3^- , pulmonary function test including FEV_1 (forced expiratory volume in 1 second; predicted), FEV_1/FVC (forced vital capacity), and pulmonary artery systolic pressure based on echocardiography were extract from the patients' files and recorded separately for each case. No PFT records were available for patients under the age of six years. Based on pulmonary artery systolic pressure (PASP), patients were then divided into two groups: 1) PH group, defined as patients with $\text{PASP} \geq 25$ mmHg and 2) non-PH group, defined as patients with $\text{PASP} < 25$ mmHg. Regarding the potential effect of age on the pulmonary artery diameter [10], the study cohort were subsequently categorised into paediatric (≤ 18 years old) and adult CF (> 18 years) groups, and all the parameters were then compared between the PH and non-PH group in each age group, using the same cut-off.

CT acquisition

CT scan was performed using a spiral CT unit (Siemens SOMATOM Emotion Series, Siemens Erlangen, Germany). All patients were imaged in the supine position, in inspiratory phases from lung apex to below the costophrenic angles, according to the standard acquisition protocol in adults and paediatric patients, as appropriate: 80-100 mA, 110 kVp, 1-1.4 pitch, and 0.5-second rotation time. CT scans were reconstructed at section widths of 1 mm and intervals of 20 mm.

CT vascular metrics

The widest short-axis diameter of the main pulmonary artery was measured on the axial section at the level of the main pulmonary artery bifurcation in mediastinal window setting (window width 400, window centre 30) on expiratory phase images. The widest short-axis diameter of ascending aorta was then measured on an axial section at the same level. The main pulmonary artery diameter (PA) to ascending aorta (A) (PA/A) ratio was then calculated. The measures were performed blinded to the haemodynamic results.

Echocardiography

Colour Doppler echocardiography was performed using Vivid 7 dimension; Mann healthcare, GE. PASP was calculated as the sum of the peak trans-tricuspid systolic pressure gradient and right atrium pressure, using modified Bernoulli equation. $\text{PASP} \geq 25$ mmHg was considered as PH.

Statistical analysis

Normally distributed data were analysed using parametric tests. Continuous (age, BMI, HCO_3 , PCO_2 , FEV_1 , FEV_1/FVC , PASP, PAD, PA/A ratio) and categorical (gender) variables were described as mean (\pm SD) and frequency (per cent), respectively. The significance of difference for each variable between two groups was tested using independent *t*-test, Mann-Whitney *U* test, and chi-square, as appropriate. Correlation between vascular and haemodynamic parameters was analysed using Pearson's correlation test. Variables with significant difference based on univariate analysis were further analysed in a logistic regression model. Diagnostic performance of pulmonary artery diameter and PA/A ratio and appropriate cut-off were calculated using receiver operating characteristic (ROC) analysis. A *p* value < 0.05 was considered as a statistically significant difference. Data were analysed using SPSS ver. 23.

Results

Of a total 119 patients with CF, 74 (62.2%) were male and 45 (38.7%) were female (*p* = 0.008), with mean age of 14.87 ± 6.4 years (age range: 3–33 years). Mean age did not demonstrate a significant difference between male and female (14.4 vs. 15.62, *p* = 0.3).

In this study the mean PAD was 19.88 mm. The mean diameter of the pulmonary artery was greater in adults with CF compared to paediatric patients (22.06 vs. 18.73, *p* = 0.000).

Regarding haemodynamic parameters, mean PASP was measured as 24.39 mmHg. Based on a PASP cut-off of 25 mmHg, 39 patients (32.8%) had increased pulmonary hypertension (mean PASP = 32.8 mmHg), and 80 (67.2%) showed normal pulmonary systolic arterial pressure (mean PASP = 20.05 mmHg). The frequency of PH and mean PASP did not show a significant difference between paediatric and adults with CF in this study (29.5% vs. 39%, *p* = 0.29 and 24.08 vs. 24.97, *p* = 0.18).

Among demographic and anthropometric characteristics, mean age was significantly higher in the PH group (16.94 vs. 13.85 years old, *p* = 0.015). The PH paediatric group had also a significantly higher mean age compared to the non-PH paediatric group (14.26 vs. 10.44 years old, *p* = 0.001); however, such a difference was not demonstrated between PH and non-PH groups in adult CF patients (22 vs. 20.81, *p* = 0.22).

Among arterial blood gas indices, mean PCO_2 and HCO_3 were found to have significantly higher values in the PH group compared to the non-PH cohort ($47.98 [\pm 9.03]$ vs. $44.11 [\pm 11.70]$, *p* = 0.011 and $29.89 [\pm 4.06]$ vs. $27.31 [\pm 6.27]$, *p* = 0.003, respectively). The same results were obtained for paediatric CF patients ($45.83 [\pm 11.43]$ vs. $43.5 [\pm 11.76]$, *p* = 0.03 and $38.92 [\pm 4.27]$ and $31.42 [\pm 3.14]$, *p* = 0.003, respectively). There was a trend toward a higher

mean HCO_3 in the CF adult PH group than in the non-PH cohort, although it was not statistically significant (*p* = 0.08) (Table 1).

There was a borderline statistical significant difference in FEV_1 between the PH and non-PH group (*p* = 0.064). While no difference was found in pulmonary function test values between the paediatric PH and non-PH group, adult PH patients had a significantly higher value of FEV_1/FVC compared to the non-PH adult cohort (*p* = 0.04).

Mean pulmonary arterial diameter and pulmonary-to-aortic (PA/A) ratio were significantly greater in the PH group than in those with normal pulmonary pressure (22.86 vs. 18.42, *p* = 0.000 and 1.03 vs. 0.95, *p* = 0.002). In the paediatric group with CF, CT vascular metrics were also significantly higher in the PH group than in the non-PH group, although such a difference was not demonstrated in adult CF patients. Table 2 summarises the CT-measured vascular parameters, based on a PASP cut-off of 25 mmHg, as the discriminator of PH from non-PH groups.

There was a statistically significant correlation between haemodynamic (PSAP) and vascular parameters (paediatric patients – PAD: [R] = 0.59, *p* = 0.000, PA/A ratio: [R] = 0.47, *p* = 0.00; adults – PAD: [R] = 0.51, *p* = 0.006, PA/A ratio: [R] = 0.45, *p* = 0.023). In paediatric patients, the correlation between PSAP and age was also significant ([R] = 0.49, *p* = 0.000). Pulmonary systolic artery pressure demonstrated a weaker inverse correlation with HCO_3 ([R] = -0.43, *p* = 0.003) and PCO_2 ([R] = -0.34, *p* = 0.029) in children.

Figure 1 demonstrates the area under the ROC curve of pulmonary artery diameter as 0.81 (CI: 0.68–0.85, *p* = 0.000) and for the PA/A ratio as 0.73 (CI: 0.57–0.78, *p* = 0.002). Using cut-off of 19.25 mm, the sensitivity and specificity of PAD for identification of PH were 82% and 70%, respectively. At a cut-off value of 0.94, the PA/A ratio demonstrated a sensitivity of 83% and specificity of 55%. The results were similar for total CF and cohort and paediatric CF patients.

Variables with a significant correlation with PSAP (age, PCO_2 , HCO_3 , PAD, PA/A ratio) based on univariate analysis were further analysed by multiple regression test, which revealed PAD as the only independent predictor of PH in total CF patients (*p* = 0.006) and in paediatric CF patients (*p* = 0.033). In adults groups, no independent relationship was found between pulmonary artery hypertension and CT-measured vascular metrics.

Discussion

The current study provided new insight into the age-dependent performance of CT-measured PAD to predict PH in CF patients. PAD may serve as a reliable tool to identify PH in paediatric patients with CF; however, no significant association was found between the level of PASP and pulmonary artery diameter in adults.

Table 1. Baseline characteristics in patients with and without pulmonary hypertension

Characteristics			Overall (n = 119)	PSAP < 25 mmHg (n = 39, 32.8%)	PSAP ≥ 25 mmHg (n = 83, 67.2%)	P value (univariate analysis)	P value (multiple logistic regression)
Demographics	Age (yr)						
		Overall	14.86 (±6.4)	13.85 (±7.08)	16.94 (±4.04)	0.003	0.97
		Pediatrics	11.36 (±4.68)	10.14 (±4.74)	14.26 (±2.78)	0.000	0.3
		Adults	21.53 (±3.03)	22 (±3.58)	20.81 (±1.76)	0.22	
	Gender						
		Overall					
	Male		74 (62.2%)	54 (72.97%)	20 (27.02%)	0.08	
	Female		45 (37.8%)	26 (57.78%)	19 (42.22%)		
		Pediatrics					
	Male		49 (62.82%)	38 (77.55%)	11 (22.45%)	0.07	
	Female		29 (31.18%)	17 (58.62%)	12 (41.39%)		
		Adults				0.62	
	Male		25 (60.97%)	16 (64%)	9 (36%)		
	Female		16 (39.02%)	7 (43.75%)	9 (56.25%)		
Anthropometrics	BMI						
		Overall					
	< 20		106 (84.1%)	72 (67.92%)	34 (32.08%)	0.6	
	20-25		13 (10.4%)	8 (61.53%)	5 (38.46%)		
		Pediatrics					
	< 20		72 (92.3%)	51 (70.8%)	21 (29.2%)	0.8	
	20-25		6 (7.7%)	4 (66.7%)	2 (33.3%)		
		Adults					
	< 20		34 (82.9%)	21 (61.8%)	13 (38.2%)	0.8	
	20-25		7 (17.1%)	4 (57.1%)	3 (42.9%)		
Arterial blood gas indices		Overall					
	pH		7.39 (±0.06)	7.4 (±0.06)	7.39 (±0.06)	0.89	
	PCo ₂ (mmHg)		45.37 (±11.01)	44.11 (±11.7)	47.99 (±9.03)	0.011	0.37
	HCO ₃ (mmol/l)		28.16 (±5.76)	27.31 (±6.29)	29.89 (±4.05)	0.003	0.15
		Pediatrics					
	pH		7.39 (±0.057)	7.4 (±0.06)	7.34 (±0.05)	0.66	
	PCo ₂ (mmHg)		44.18 (±11.64)	43.5 (±11.76)	45.83 (±11.43)	0.03	0.64
	HCO ₃ (mmol/l)		33.63 (±35.96)	34.42 (±32.91)	38.93 (±42.72)	0.003	0.5
		Adults					
	pH		7.39 (±0.07)	7.34 (±0.06)	7.39 (±0.08)	0.82	
	PCo ₂ (mmHg)		45.91 (±12.2)	44.08 (±13.46)	48.78 (±4.64)	0.32	
	HCO ₃ (mmol/l)		34.18 (±3.8)	27.09 (±6.7)	45.26 (±5.97)	0.08	
Pulmonary function test		Overall					
	FEV ₁		53.47 (±25.69)	57.14 (±26.26)	47.57 (±23.84)	0.064	
	FEV ₁ /FVC		97.12 (±14.73)	94.14 (±15.98)	98.79 (±12.33)	0.61	
		Pediatric					
	FEV ₁		56.91 (±28.23)	55.28 (±31.86)	46.3 (±26.11)	0.29	
	FEV ₁ /FVC		97.03 (±16.33)	97.33 (±17.96)	96.47 (±13.2)	0.56	
		Adults					
	FEV ₁		48.63 (±20.97)	49.32 (±21.02)	47.36 (±21.52)	0.55	
	FEV ₁ /FVC		97.26 (±11.95)	94.16 (±12.00)	102.12 (±10.44)	0.04	

Table 2. Pulmonary hemodynamic and vascular metrics in patients with and without pulmonary hypertension

Characteristics		Overall (n = 119)	PSAP < 25 mmHg (n = 83, 67.2%)	PSAP ≥ 25 mmHg (n = 39, 32.8%)	P value (univariate analysis)	P value (multiple logistic regression)
Pulmonary systolic pressure (mmHg)						
	Overall	24.39 (±8.85)	20.05 (±7.93)	33.30 (±10.98)	0.00	
	Pediatrics	24.08 (±8.46)	19.94 (±7.3)	34 (±10.69)	0.00	
	Adults	24.97 (±9.32)	20.28 (±8.9)	32.31 (±11.66)	0.000	
Pulmonary artery diameter (mm)						
	Overall	19.88 (±4.91)	18.43 (±4.72)	22.86 (±3.86)	0.000	0.005
	Pediatrics	18.73 (±5.13)	17.10 (±4.64)	22.63 (±4.4)	0.000	0.03
	Adults	22.06 (±3.61)	21.34 (±3.49)	23.19 (±3.60)	0.7	
PA/A						
	Overall	0.98 (±0.12)	0.95 (±0.11)	1.03 (±0.13)	0.002	0.79
	Pediatrics	0.99 (±0.12)	0.97 (±0.11)	1.05 (±0.13)	0.001	0.77
	Adults	0.96 (±0.13)	0.94 (±0.12)	1.00 (±0.14)	0.125	

Few studies are available investigating various aspects of PH in CF. It has been shown that PH is frequently seen in advanced CF (57%) [10]. In the present study, the frequency of PH was found to be 32.8%. This difference may be at least partly due to the lower prevalence of advanced CF in the current study (2.5%).

Pulmonary hypertension is a pathophysiological condition caused by a variety of diseases mediated via different underlying anatomical and physiological mechanisms. In chronic obstructive pulmonary disease (COPD), recent studies have demonstrated that a direct toxic effect of tobacco on the intima proceed vascular remodelling via abnormal production of the mediators controlling vasodilation and vasoconstriction and also abnormal intimal cell proliferation [11,12]; however, in CF patients, vascular remodelling is strongly influenced by the alveolar hypoxia as well as a chronic systemic inflammatory condition [13,14]. Augmentation index (AIx), as a significant predictor of arterial stiffness with a potential association with systemic inflammation, has been demonstrated to have higher values in the adult CF population, indicating an accelerated vascular ageing process in this cohort [14]. Despite an independent relationship in paediatric CF, the present study failed to demonstrate any striking correlation between pulmonary artery diameter and PH in adult CF. Taking this into consideration, one possible explanation is the well-established premature vascular ageing and advanced arterial stiffness in adult CF compared to the paediatric group, preventing a linear relationship between pulmonary artery pressure and pulmonary artery diameter.

Most researches demonstrated that pulmonary haemodynamic indices are independent of lung function status in CF patients [13], which is in accordance with the results obtained in the present study. Such results pro-

vided evidence for the major limitation of PFT to be solely applied for disease monitoring in CF patients, and they support the potential value of CT-measured thoracic vascular diameter as a complementary test to PFT for comprehensive disease monitoring in paediatric CF patients.

Several reports revealed a significant correlation between obesity and PH. One study demonstrated that 72% of subjects with BMI > 30 kg/m² showed haemodynamic changes in pulmonary circulation, including venous hypertension and alveolar capillary haemangiomatosis, indicating of an underlying PH [15]. Another study demonstrated that the frequency of obesity was significantly higher in patients with PH than in those with normal pulmonary pressure [16]. The same result was also observed in COPD patients with pulmonary hypertension – mean BMI was significantly higher in patients with PH (28 vs. 25, $p = 0.04$) [17]. However, the current study failed to provide any association between BMI and pulmonary systolic arterial pressure as well as CT vascular metrics. This discrepancy could be explained by the confounding effect of underlying malnutrition status most frequently seen in CF patients as a result of pulmonary function disturbance and gastrointestinal complications including pancreatic insufficiency, steatorrhoea, hepatic steatosis, and biliary fibrosis.

The present study has some major drawbacks. Retrospective data collection from a single CF centre may limit the validity of results to the CF patient population. However, it should be noted that Masih Daneshvari Hospital, as a tertiary referral centre for CF with a comprehensive CF registry, provides a wide coverage of CF patients in Iran. The impact of CT vascular metrics to predict PH on clinical outcome needs to be further investigated. None of the patients were candidates for right-sided heart catheterisation, and hence the exact value of haemodynamic parameters measured by gold standard were not available;

however, the current results may be less influenced by the selection bias in the studies performed on RVH candidate cohorts. Associated pulmonary parenchymal involvement as a major determinant of survival with a potential impact on pulmonary arterial systolic pressure was not evaluated in this study. Small subsets of patients in age groups and limited age range of adult CF (up to 33 years old) should be considered as the important consideration for the present results.

In conclusion, the current study provided evidence for the age-dependent performance of CT-based pulmonary artery diameter, as an easily obtainable measurement, to predict PH. PAD correlates well with PH in paediatric

patients with CF, and this relationship is independent of age, lung function test, and arterial blood gas indices. In this regard, CT-measured thoracic vascular diameter may be recommended as a complementary test to pulmonary function test for monitoring of disease progression in childhood CF. However, the added value of CT-based follow-up approach in paediatric CF should be further validated by large-scale prospective survival studies considering excess radiation exposure.

Conflict of interest

The authors declare that they have no conflict of interest.

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